## DEPARTMENT OF JUSTICE

**Drug Enforcement Administration** 

[Docket No. DEA-392]

Manufacturer of Controlled Substances Registration: Austin Pharma LLC

**ACTION:** Notice of registration.

**SUMMARY:** Austin Pharma LLC applied to be registered as a manufacturer of certain basic classes of controlled substances. The Drug Enforcement Administration (DEA) grants Austin Pharma LLC registration as a manufacturer of those controlled substances.

## **SUPPLEMENTARY INFORMATION:**

By notice dated August 10, 2015, and published in the *Federal Register* on August 18, 2015, 80 FR 50043, Austin Pharma LLC, 811 Paloma Drive, Suite C, Round Rock, Texas 78665-2402 applied to be registered as a manufacturer of certain basic classes of controlled substances. No comments or objections were submitted for this notice.

The DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Austin Pharma LLC to manufacture the basic classes of controlled substances is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. The DEA investigated the company's maintenance of effective controls against diversion by inspecting and testing the company's physical security systems, verifying the company's compliance with state and local laws, and reviewing the company's background and history.

Therefore, pursuant to 21 U.S.C. 823(a), and in accordance with 21 CFR 1301.33, the above-named company is granted registration as a bulk manufacturer of the following basic classes of controlled substances:

Controlled Substance	<u>Schedule</u>
Marihuana (7360)	Ι
Tetrahydrocannabinols (7370)	I

The company plans to manufacture bulk synthetic active pharmaceutical ingredients (APIs) for product development and distribution to its customers. No other activity for these drug codes are authorized for this registration.

Dated: December 9, 2015.

Louis J. Milione, *Deputy Assistant Administrator*.

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